Amendments to the Claims

1-6. (Cancelled)

7. (Currently Amended) Isolated and purified biologically active TFPI comprising an N-terminal amino acid sequence as shown in SEQ ID NO: 7, wherein the biologically active TFPI has an inhibitory concentration of at least 1 µg/ml in a prothrombin clotting assay, according to made by a method comprising:

transforming yeast cells with a vehicle, said vehicle comprising a first nucleotide sequence encoding a first protein, wherein the first protein is TFPI, wherein the N-terminal amino acid sequence of the TFPI is SEQ ID NO: 7, said first nucleotide sequence being immediately preceded in frame by a second nucleotide sequence encoding a second protein ubiquitin, the first and second nucleotide sequences together encoding a fusion protein;

incubating the transformed yeast cells under conditions favorable for production of the whereby the fusion protein is produced and cleaved to produce TFPI, wherein the TFPI is retained within the yeast cell;

preparing an insoluble fraction of the transformed yeast cells containing the TFPI; and recovering the TFPI from the insoluble fraction.

wherein the TFPI comprises the N-terminal amino acid sequence shown in SEQ ID NO:7 and wherein the TFPI has an inhibitory concentration of at least 1 μg/ml in a prothrombin clotting assay.

8-11. (Cancelled)

(New) The isolated and purified TFPI of claim 7 wherein the yeast cell of the method is a

Saccharomyces cerevisiae cell having a genotype selected from the group consisting of VH6,

AB122, and JSC310.

- 13. (New) The method of claim 7 wherein the second protein is ubiquitin.
- 14. (New) The method of claim 7 wherein the second protein is superoxide dismutase.